

L Number	Hits	Search Text	DB	Time stamp
1	404	metronidazole	EPO; JPO; DERWENT	2003/01/03 10:31
2	7273	cyclodextrin	EPO; JPO; DERWENT	2003/01/03 10:31
3	504	niacin niacinamide	EPO; JPO; DERWENT	2003/01/03 10:31
4	5318	nicotinic nicotinamide	EPO; JPO; DERWENT	2003/01/03 10:31
5	2008371	water aqueous solubil\$9	EPO; JPO; DERWENT	2003/01/03 10:32
6	128	(cyclodextrin (niacin niacinamide) (nicotinic nicotinamide) (water aqueous solubil\$9)) and metronidazole	EPO; JPO; DERWENT	2003/01/03 10:32

L Number	Hits	Search Text	DB	Time stamp
1	1870	metronidazole	USPAT; US-PGPUB	2003/01/03 09:38
2	1056608	aqueous water solubil\$9	USPAT; US-PGPUB	2003/01/03 09:39
3	187	metronidazole same (aqueous water solubil\$9)	USPAT; US-PGPUB	2003/01/03 09:39
4	7770	cyclodextrin	USPAT; US-PGPUB	2003/01/03 09:39
5	13271	nicotinic nicotinamide	USPAT; US-PGPUB	2003/01/03 09:39
6	3623	niacin niacinamide	USPAT; US-PGPUB	2003/01/03 09:40
7	24505	cyclodextrin (nicotinic nicotinamide) (niacin niacinamide) metronidazole	USPAT; US-PGPUB	2003/01/03 09:40
8	187	(metronidazole same (aqueous water solubil\$9)) and (cyclodextrin (nicotinic nicotinamide) (niacin niacinamide) metronidazole)	USPAT; US-PGPUB	2003/01/03 09:40

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(FILE 'HOME' ENTERED AT 08:52:38 ON 03 JAN 2003)

FILE 'REGISTRY' ENTERED AT 08:52:50 ON 03 JAN 2003

L1       1 S METRONIDAZOLE/CN  
          SELECT L1 1- CHEM

FILE 'CAPLUS, MEDLINE, EMBASE, BIOSIS' ENTERED AT 08:53:58 ON 03 JAN 2003

L2       47618 S E1-32  
L3       32748 DUP REM L2 (14870 DUPLICATES REMOVED)  
L4       40689 S CYCLODEXTRIN  
L5       86817 S NICOTINIC  
L6       67402 S NICOTINAMIDE  
L7       11231 S NIACIN  
L8       198140 S L4 OR L5 OR L6 OR L7  
L9       46716 S METRONIDAZOLE  
L10      40689 S CYCLODEXTRIN  
L11      86817 S NICOTINIC  
L12      67402 S NICOTINAMIDE  
L13      11231 S NIACIN  
L14      5763 S NIACINAMIDE  
L15      201186 S L10 OR L11 OR L12 OR L13 OR L14  
L16      189 S L15 AND L9  
L17      161 DUP REM L16 (28 DUPLICATES REMOVED)  
L18      435595 S SOLUBILITY OR SOLUBILIZ?  
L19      1161421 S AQUEOUS  
L20      3097083 S WATER  
L21      3932107 S L19 OR L20  
L22      13 S L17 AND L18 AND L21

L22 ANSWER 1 OF 13 CAPLUS COPYRIGHT 2003 ACS  
 ACCESSION NUMBER: 2002:72162 CAPLUS  
 DOCUMENT NUMBER: 136:107569  
 TITLE: Gel compositions containing metronidazole  
 and hydroxypropyl-.beta.-cyclodextrin  
 INVENTOR(S): Chang, Yunik; Dow, Gordon J.; Angel, Arturo  
 PATENT ASSIGNEE(S): Dow Pharmaceutical Sciences, USA  
 SOURCE: PCT Int. Appl., 35 pp.  
 CODEN: PIXXD2  
 DOCUMENT TYPE: Patent  
 LANGUAGE: English  
 FAMILY ACC. NUM. COUNT: 1  
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2002006349	A1	20020124	WO 2001-US19644	20010619
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG				
US 6468989	B1	20021022	US 2000-615169	20000713

PRIORITY APPLN. INFO.: US 2000-615169 A 20000713

AB An aq. soln. of metronidazole in which the concn. of metronidazole is >0.75 is described. The soln. contains the solv. enhancer hydroxypropyl-.beta.-cyclodextrin (I) and may addnl. contain niacinamide. Methods of manuf. and therapeutic use of the soln. are disclosed. Thus, a stable 1.0% aq. gel compn. contained metronidazole 1.00, I 5.00, methylparaben 0.15, propylparaben 0.03, glycerin 5.00, hydroxyethyl cellulose 1.50, disodium edetate 0.05, and water qs to 100%.

REFERENCE COUNT: 3 THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L22 ANSWER 2 OF 13 CAPLUS COPYRIGHT 2003 ACS  
 ACCESSION NUMBER: 2002:71873 CAPLUS  
 DOCUMENT NUMBER: 136:123671  
 TITLE: Ophthalmic formulation of a selective cyclooxygenase-2 inhibitory drug  
 INVENTOR(S): Kararli, Tugrul T.; Bandyopadhyay, Rebanta; Singh, Satish K.; Hawley, Leslie C.  
 PATENT ASSIGNEE(S): Pharmacia & Upjohn Company, USA  
 SOURCE: PCT Int. Appl., 71 pp.  
 CODEN: PIXXD2  
 DOCUMENT TYPE: Patent  
 LANGUAGE: English  
 FAMILY ACC. NUM. COUNT: 4  
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2002005815	A1	20020124	WO 2001-US22061	20010712
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG				
US 2002035264	A1	20020321	US 2001-904098	20010712
PRIORITY APPLN. INFO.:			US 2000-218101P	P 20000713
			US 2001-279285P	P 20010328
			US 2001-294838P	P 20010531
			US 2001-296388P	P 20010606

OTHER SOURCE(S): MARPAT 136:123671

AB A pharmaceutical compn. suitable for topical administration to an eye contains a selective COX-2 inhibitor or nanoparticles of a drug of low water solv., at a concn. effective for the treatment and/or prophylaxis of a disorder in the eye, and 1 or more ophthalmically acceptable excipients that reduce rate of removal from the eye such that

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the compn. has an effective residence time of 2-24 h. Also provided is a method of treating and/or preventing a disorder in an eye, the method comprising administering to the eye a compn. of the invention. Thus, an ophthalmic nanoparticle suspension contained valdecoxib at 2.15 mg/g, 1.2% glycerin, 0.8% EDTA disodium salt, 4.0% Gelcarin GP-379NF, 0.21% SeaSpen PF and 0.82% Povidone.

REFERENCE COUNT: 10 THERE ARE 10 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L22 ANSWER 3 OF 13 CAPLUS COPYRIGHT 2003 ACS

ACCESSION NUMBER: 2001:300514 CAPLUS

DOCUMENT NUMBER: 134:331617

TITLE: Oil-in-water emulsion compositions for polyfunctional active ingredients

INVENTOR(S): Chen, Feng-jing; Patel, Mahesh V.

PATENT ASSIGNEE(S): Lipocene, Inc., USA

SOURCE: PCT Int. Appl., 82 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2001028555	A1	20010426	WO 2000-US28835	20001018
W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM			
RW:	GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG			
US 2002107265	A1	20020808	US 1999-420159	19991018

PRIORITY APPLN. INFO.: US 1999-420159 A 19991018

AB Pharmaceutical oil-in-water emulsions for delivery of polyfunctional active ingredients with improved loading capacity, enhanced stability, and reduced irritation and local toxicity are described. Emulsions include an aq. phase, an oil phase comprising a structured triglyceride, and an emulsifier. The structured triglyceride of the oil phase is substantially free of triglycerides having three medium chain (C6-C12) fatty acid moieties, or a combination of a long chain triglyceride and a polarity-enhancing polarity modifier. The present invention also provides methods of treating an animal with a polyfunctional active ingredient, using dosage forms of the pharmaceutical emulsions. For example, an emulsion was prep'd., with cyclosporin A as the polyfunctional active ingredient dissolved in an oil phase including a structured triglyceride (Captex 810D) and a long chain triglyceride (safflower oil). The compn. contained (by wt.) cyclosporin A 1.0, Captex 810D 5.0, safflower oil 5.0, BHT 0.02, egg phospholipid 2.4, dimyristoylphosphatidyl glycerol 0.2, glycerol 2.25, EDTA 0.01, and water up to 100%, resp.

REFERENCE COUNT: 6 THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L22 ANSWER 4 OF 13 CAPLUS COPYRIGHT 2003 ACS

ACCESSION NUMBER: 1996:71430 CAPLUS

DOCUMENT NUMBER: 124:155977

TITLE: Cyclodextrin complexation

INVENTOR(S): Loftsson, Thorsteinn

PATENT ASSIGNEE(S): Cyclops h.f., Iceland

SOURCE: U.S., 31 pp. Cont.-in-part of U.S. 5,324,718.

CODEN: USXXAM

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 2

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 5472954	A	19951205	US 1994-240510	19940511
US 5324718	A	19940628	US 1992-912853	19920714
EP 579435	A1	19940119	EP 1993-305280	19930706
EP 579435	B1	19990317		

R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LI, LU, MC, NL, PT, SE  
PRIORITY APPLN. INFO.: US 1992-912853 19920714

EP 1993-305280 19930706

AB The invention provides a method for enhancing the complexation of a cyclodextrin with a lipophilic and/or water-labile active ingredient which is a drug, cosmetic additive, food additive or agrochem., comprising combining from about 0.1 to about 70% (wt./vol.) of a cyclodextrin, from about 0.001 to about 5% (wt./vol.) of a pharmacol. inactive water-sol. polymer acceptable for use in a pharmaceutical, cosmetic, food or agricultural compn., and said lipophilic and/or water-labile active ingredient in an aq. medium, the polymer and cyclodextrin being dissolved in the aq. medium before the active ingredient is added, the aq. medium which comprises the polymer and cyclodextrin being maintained at 30-150.degree. for 0.1-100 h before, during and/or after the active ingredient is added, optionally followed by removal of water. Related methods, co-complexes of active ingredient/cyclodextrin/polymer, pharmaceutical, cosmetic, food and agricultural compns. and cyclodextrin/polymer complexing agents are also provided.

L22 ANSWER 5 OF 13 CAPLUS COPYRIGHT 2003 ACS

ACCESSION NUMBER: 1995:454268 CAPLUS

DOCUMENT NUMBER: 122:298838

TITLE: Preparation and characterization of metronidazole benzoate-.gamma.-cyclodextrin inclusion compound

AUTHOR(S): Giordano, F.; Bruni, G.; Abdel Hadi, Ismail; Kata, Mihaly; Gazzaniga, A.; Bettinetti, G.

CORPORATE SOURCE: Dipartimento di Chimica Farmaceutica, Univ. di Pavia, Pavia, 27100, Italy

SOURCE: Bollettino Chimico Farmaceutico (1992); 131(4), 150-6  
CODEN: BCFAAI; ISSN: 0006-6648

PUBLISHER: Societa Editoriale Farmaceutica

DOCUMENT TYPE: Journal

LANGUAGE: English

AB Metronidazole benzoate, an antiprotozoal drug, forms a 1:1 (mol/mol) inclusion compd. with .gamma.-cyclodextrin. Phase-solv. anal., differential scanning calorimetry, x-ray diffraction on powder, and IR spectra were used in order to characterize the inclusion compd. both in soln. and in solid state. The stability of the drug to alk. hydrolysis was improved in aq. solns. of .gamma.-cyclodextrin.

L22 ANSWER 6 OF 13 CAPLUS COPYRIGHT 2003 ACS

ACCESSION NUMBER: 1994:517696 CAPLUS

DOCUMENT NUMBER: 121:117696

TITLE: Derivatives of cyclodextrins exhibiting enhanced aqueous solubility and the use thereof

INVENTOR(S): Stella, Valentino J.; Rajewski, Roger

PATENT ASSIGNEE(S): University of Kansas, USA

SOURCE: PCT Int. Appl., 72 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 3

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 9402518	A1	19940203	WO 1993-US6880	19930726
W: AU, CA, JP, KR, RU				
RW: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE				
JP 06511513	T2	19941222	JP 1992-504678	19920726
US 5376645	A	19941227	US 1992-918702	19920727
EP 620828	A1	19941026	EP 1993-918302	19930726
EP 620828	B1	20020508		
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LI, LU, MC, NL, PT, SE				
AU 672814	B2	19961017	AU 1993-47799	19930726
AU 9347799	A1	19940214		
AT 217325	E	20020515	AT 1993-918302	19930726
PRIORITY APPLN. INFO.:			US 1992-918702	A 19920727
			US 1990-469087	A2 19900123
			WO 1993-US6880	W 19930726

OTHER SOURCE(S): MARPAT 121:117696

AB Sulfoalkyl ether cyclodextrin derivs. and their use as solubilizing agents for water insol. drugs for oral, intranasal, or parenteral administration are disclosed. For example, .beta.-cyclodextrin sulfopropyl ether (7 substituents per

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cyclodextrin mol.) was prep'd. and assocn. consts. for the equil. between the sulfopropyl derivs. and drugs, i.e. digoxin, progesterone, testosterone, and phenytoin were studied.

L22 ANSWER 7 OF 13 CAPLUS COPYRIGHT 2003 ACS

ACCESSION NUMBER: 1994:253358 CAPLUS

DOCUMENT NUMBER: 120:253358

TITLE: Cyclodextrin complexes with polymers, drugs, agrochemicals and cosmetics

INVENTOR(S): Loftsson, Thorsteinn

PATENT ASSIGNEE(S): Iceland

SOURCE: Eur. Pat. Appl., 46 pp.

CODEN: EPXXDW

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 2

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
EP 579435	A1	19940119	EP 1993-305280	19930706
EP 579435	B1	19990317		
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LI, LU, MC, NL, PT, SE			
US 5324718	A	19940628	US 1992-912853	19920714
AT 177647	E	19990415	AT 1993-305280	19930706
ES 2132190	T3	19990816	ES 1993-305280	19930706
US 5472954	A	19951205	US 1994-240510	19940511
PRIORITY APPLN. INFO.:		US 1992-912853	19920714	
		EP 1993-305280	19930706	

AB A method for enhancing the complexation of a cyclodextrin (I) with a lipophilic and/or water-labile drug, comprising combining .apprx.0.1-70% (wt./vol.) of I and .apprx.0.001-5% (wt./vol.) of a water-sol. polymer in an aq. medium. The polymer and I are dissolved in the aq. medium before the drug is added. To a soln. contg. Na CM-cellulose 0.25 and 2-hydroxypropyl-.beta.-cyclodextrin 10% was added acetazolamide (II) and the soln. was heated at 120.degree. for 20 min and allowed to equilibrate at room temp. for 3 days and amt. of II was detd. The solv. of II was 3.11mg/mL as compared to 0.7 for control contg. only II. Different formulations contg. cyclodextrin complexes with polymers and drugs are disclosed.

L22 ANSWER 8 OF 13 CAPLUS COPYRIGHT 2003 ACS

ACCESSION NUMBER: 1991:589787 CAPLUS

DOCUMENT NUMBER: 115:189787

TITLE: Derivatives of cyclodextrins exhibiting enhanced aqueous solubility and the use thereof

INVENTOR(S): Stella, Valentino; Rajewski, Roger

PATENT ASSIGNEE(S): University of Kansas, USA

SOURCE: PCT Int. Appl., 48 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 3

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 9111172	A1	19910808	WO 1991-US326	19910122
	W: AU, CA, JP, KR, SU			
	RW: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LU, NL, SE			
US 5134127	A	19920728	US 1990-469087	19900123
CA 2074186	AA	19910724	CA 1991-2074186	19910122
AU 9172364	A1	19910821	AU 1991-72364	19910122
AU 646020	B2	19940203		
EP 512050	A1	19921111	EP 1991-903891	19910122
EP 512050	B1	19980909		
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE			
JP 05504783	T2	19930722	JP 1991-504051	19910122
JP 2722277	B2	19980304		
AT 170742	E	19980915	AT 1991-903891	19910122
RU 2099354	C1	19971220	RU 1992-5052811	19920722
PRIORITY APPLN. INFO.:		US 1990-469087	A 19900123	
		WO 1991-US326	A 19910122	

OTHER SOURCE(S): MARPAT 115:189787

AB Cyclodextrin sulfoalkyl ethers (Markush given) are prep'd. as clathrating agents to enhance the water solv. of

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drugs. A mixt. contg. .beta.-cyclodextrin 5, NaOH 2 g, and 10 mL water was treated with 4.5 mL of butane sultone and the resulting soln. was neutralized with 1 N HCl to give sulfobutyl ether of .beta.-cyclodextrin. The product exhibited no observable toxic effects in mice over a 30 day period following i.p. injection of 0.00549 mol/kg.

L22 ANSWER 9 OF 13 CAPLUS COPYRIGHT 2003 ACS  
ACCESSION NUMBER: 1990:446267 CAPLUS  
DOCUMENT NUMBER: 113:46267  
TITLE: Pharmaceutical formulations for parenteral use containing cyclodextrins and dihydropyridine redox systems  
INVENTOR(S): Bodor, Nicholas S.  
PATENT ASSIGNEE(S): University of Florida, USA  
SOURCE: Eur. Pat. Appl., 125 pp.  
CODEN: EPXXDW  
DOCUMENT TYPE: Patent  
LANGUAGE: English  
FAMILY ACC. NUM. COUNT: 2  
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
EP 335545	A2	19891004	EP 1989-302719	19890320
EP 335545	A3	19900926		
EP 335545	B1	19930609		
EP 335545	B2	19980923		
R: AT, BE, CH, DE, ES, FR, GB, GR, IT, LI, LU, NL, SE				
US 4983586	A	19910108	US 1988-174945	19880329
EP 327766	A2	19890816	EP 1988-312016	19881219
EP 327766	A3	19900926		
EP 327766	B1	19980408		
R: AT, BE, CH, DE, ES, FR, GB, GR, IT, LI, LU, NL, SE				
AT 90200	E	19930615	AT 1989-302719	19890320
AU 8931762	A1	19890727	AU 1989-31762	19890328
AU 618995	B2	19920116		
CA 1336498	A1	19950801	CA 1989-594911	19890328
JP 02009825	A2	19900112	JP 1989-77938	19890329
JP 2643426	B2	19970820		
ZA 8902315	A	19901228	ZA 1989-2315	19890329
US 5017566	A	19910521	US 1989-431222	19891103
US 5024998	A	19910618	US 1989-448655	19891211
PRIORITY APPLN. INFO.:			US 1988-174945	A 19880329
			EP 1988-312016	A 19881219
			US 1987-139755	A2 19871230
			CA 1988-585791	A 19881213
			IE 1988-3717	A 19881213
			IE 1989-810	A 19890314
			EP 1989-302719	A 19890320
			US 1989-431222	A2 19891103

AB Aq. parenteral solns. of drugs which are insol. or only sparingly sol. and/or which are unstable in water, are combined with a cyclodextrin deriv. to provide a means for alleviating problems assocd. with drug pptn. at the injection site and/or in the lungs or other organs following parenteral administration. Another approach is use of the dihydropyridine-pyridinium redox delivery system. A large no. of examples are given for synthesis of dihydropyridine and pyridinium derivs. of drugs. Data are also presented showing drug solubilization by cyclodextrin derivs.

L22 ANSWER 10 OF 13 CAPLUS COPYRIGHT 2003 ACS  
ACCESSION NUMBER: 1984:536904 CAPLUS  
DOCUMENT NUMBER: 101:136904  
TITLE: Inclusion complexation of metronidazole benzoate with .beta.-cyclodextrin and its depression of anhydrate-hydrate transition in aqueous suspensions  
AUTHOR(S): Andersen, Finn M.; Bundgaard, Hans  
CORPORATE SOURCE: Dep. Pharm. Chem., R. Dan. Sch. Pharm., Copenhagen, DK-2100, Den.  
SOURCE: International Journal of Pharmaceutics (1984), 19(2), 189-97  
CODEN: IJPHDE; ISSN: 0378-5173  
DOCUMENT TYPE: Journal  
LANGUAGE: English  
AB Metronidazole benzoate (I) [13182-89-3] formed an inclusion complex with .beta.-cyclodextrin (.beta.-CyD) in aq.

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soln. and in the solid phase. A phase solv. diagram was obtained and an apparent 1:1 formation complex const. of 1.3 times. 103 M-1 was detd. A microcryst. inclusion complex had the stoichiometric compn. of 1:1.5 (drug-.beta.-CyD). By inclusion complexation of the I with .beta.-CyD the phase transition of the clin. used anhyd. form of the compd. to the monohydrate, occurring in aq. suspensions, was inhibited as was the marked crystal growth resulting from the phase transition. Besides increasing the phys. stability of I suspensions, complexation with .beta.-CyD protected the drug against photochem. degrdn. and decreased the rate of hydrolysis.

L22 ANSWER 11 OF 13 CAPLUS COPYRIGHT 2003 ACS

ACCESSION NUMBER: 1984:180054 CAPLUS

DOCUMENT NUMBER: 100:180054

TITLE: Solubilization of metronidazole by water-miscible multi-cosolvents and water-soluble vitamins

AUTHOR(S): Chien, Yie W.

CORPORATE SOURCE: Coll. Pharm., Rutgers Univ., Piscataway, NJ, USA

SOURCE: Journal of Parenteral Science and Technology (1984), 38(1), 32-6

CODEN: JPATDS; ISSN: 0279-7976

DOCUMENT TYPE: Journal  
LANGUAGE: English

AB In the systemic treatment of anaerobic infections, parenteral administration of metronidazole (I) [443-48-1] is preferable. The practical way of administering a parenteral dose (500 mg) of I in a single 10-mL form can be achieved by incorporating .gtoreq.2 water-miscible cosolvents, e.g., ethanol [64-17-5], N,N-dimethylacetamide [127-19-5], propylene glycol [57-55-6], or solketal [100-79-8], into the aq. soln. The aq. solv. of I increased exponentially with increasing vol. fraction of the cosolvents. A max. solv. of I was obsd. in aq. solns. with a dielec. const. of 41.49. The importance of dielec. const. in detg. the aq. solv. of slightly water-sol. I, which consists of lipophilic and hydrophilic functional groups, is discussed. The aq. solv. of I can also be enhanced by solubilizing with a water-sol. vitamin, e.g., nicotinamide [98-92-0], ascorbic acid [50-81-7], or pyridoxine-HCl [58-56-0]. A cage-like structure was postulated to rationalize the observation that 9 mols. of vitamins are required to solubilize every mol. of I in the aq. soln.

L22 ANSWER 12 OF 13 EMBASE COPYRIGHT 2003 ELSEVIER SCI. B.V.

ACCESSION NUMBER: 95257734 EMBASE

DOCUMENT NUMBER: 1995257734

TITLE: Bioactivation of dinitrobenzamide mustards by an E. Coli B nitroreductase.

AUTHOR: Anlezark G.M.; Melton R.G.; Sherwood R.F.; Wilson W.R.; Denny W.A.; Palmer B.D.; Knox R.J.; Friedlos F.; Williams A.

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SUMMARY LANGUAGE: English

AB A nitroreductase isolated and purified from Escherichia coli B has been demonstrated to have potential applications in ADEPT (antibody-directed enzyme prodrug therapy) by its ability in vitro to reduce dinitrobenzamides (e.g. 5-aziridinyl 2,4-dinitrobenzamide, CB 1954 and its bischloroethylamino analogue, SN 23862) to form cytotoxic derivatives. In contrast to CB 1954, in which either nitro group is reducible to the corresponding hydroxylamine, SN 23862 is reduced by the nitroreductase to form only the 2-hydroxylamine. This hydroxylamine can react with S-acetylthiocholine to form a species capable of producing interstrand crosslinks in naked DNA. In terms of ADEPT, SN 23862 has a potential advantage over CB 1954 in that it is not reduced by mammalian DT diaphorases. Therefore, a series of compounds related to SN 23862 has been synthesized, and evaluated as potential prodrugs both by determination of kinetic parameters and by ratio of IC50 against UV4 cells when incubated

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in the presence of prodrug, with and without the E. coli enzyme and cofactor (NADH). Results from the two studies were generally in good agreement in that compounds showing no increase in cytotoxicity in presence of enzyme and cofactor were not substrates for the enzyme. None of the analogues were activated by DT diaphorase isolated from Walker 256 carcinoma cells. For those compounds which were substrates for the E. coli nitroreductase, there was a positive correlation was between k(cat) and IC<sub>50</sub> ratio. Two compounds showed advantageous properties: SN 25261 (with a dihydroxypropylcarboxamide ring substituent) which has a more than 10-fold greater aqueous solubility than SN 23862 whilst retaining similar kinetic characteristics and where a change in the position of the carboxamide group relative to the cytotoxicity ratio and k(cat) compared with SN 23862 (IC<sub>50</sub> ratios 214 26.4 sec-1, respectively). An analogue (SN 25507) incorporating both enhanced k(cat) of 576 sec-1. This study elucidates some of the structural aids identification of further directions in the search for suitable prodrugs system.

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TITLE: Gel compositions containing metronidazole.

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AB An aqueous solution of metronidazole in which the concentration of metronidazole is higher than 0.75%. The solution contains the solubility enhancer hydroxypropyl-beta-cyclodextrin and may additionally contain niacinamide.

Methods of manufacture and therapeutic use of the solution are disclosed.